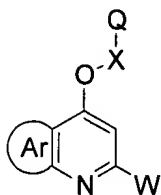
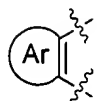


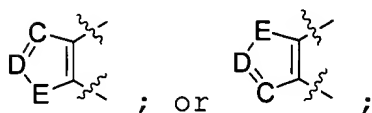
1. (Previously Presented) A compound of the formula:



or a pharmaceutically acceptable salt thereof, wherein:



represents:



wherein:

C and D are CR_1 , and

E represents sulfur,

where

R_1 , at each occurrence, is independently selected from the group consisting of hydrogen, halogen, cyano, halo(C_{1-6})alkyl, halo(C_{1-6})alkoxy, hydroxy, C_{1-6} alkyl, amino, mono and di(C_{1-6})alkylamino, and C_{1-6} alkoxy; and

R_2 is selected from the group consisting of hydrogen, halogen, cyano, halo($\text{C}_1\text{-C}_6$)alkyl, halo($\text{C}_1\text{-C}_6$)alkoxy, hydroxy, C_{1-6} alkyl, amino, and mono or di($\text{C}_1\text{-C}_6$)alkylamino;

W is phenyl which is unsubstituted or substituted with 1, 2, 3, 4, or 5 R_3 groups or naphthyl which is unsubstituted or substituted with 1, 2, 3, 4, 5, 6, or 7 R_3 groups; and

Q is pyridinyl, which is unsubstituted or substituted with 1, 2, 3, or 4 R_4 groups;

R_3 and R_4 at each occurrence are independently selected from the group consisting of hydrogen, halogen, hydroxy, $-OR_6$, $-NO_2$, $-CN$, $-SO_2NH_2$, $-SO_2NHR_6$, $-SO_2N(R_6)_2$, amino, $-NHR_6$, $-N(R_6)_2$, $-N(R_6)CO(R_6)$, $-N(R_6)CO_2(R_6)$, $-CONH_2$, $-CONH(R_6)$, $-CON(R_6)_2$, $-CO_2(R_6)$, $-S(R_6)$, $-SO(R_6)$, $-SO_2(R_6)$, and R_7 , wherein

R_6 , at each occurrence, is independently C_{1-8} alkyl, which is unsubstituted or substituted with one or two substituents independently selected from the group consisting of hydroxy, oxo, halogen, amino, and C_{1-8} alkoxy,

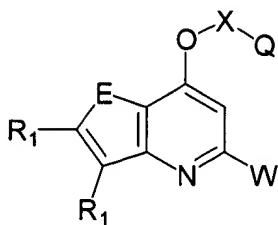
R_7 at each occurrence is independently C_{1-8} alkyl, which is unsubstituted or substituted with one or two substituents independently selected from the group consisting of

hydroxy, oxo, halogen, $-OR_6$, $-NO_2$, $-CN$, $-SO_2NH_2$, $-SO_2NHR_6$, $-SO_2N(R_6)_2$, amino, $-NHR_6$, $-N(R_6)_2$, $-N(R_6)CO(R_6)$, $-N(R_6)CO_2(R_6)$, $-CONH_2$, $-CONH(R_6)$, $-CON(R_6)_2$, $-CO_2H$, $-CO_2(R_6)$, $-S(R_6)$, $-SO(R_6)$, and $-SO_2(R_6)$,

X is $-(CH_2)_n-$ or $-(CH_2)_n(C=O)-$, wherein each n is independently 1, 2, or 3.

2-8. (Cancelled)

9. (Original) A compound or salt according to claim 1 of formula:



10. (Cancelled)

11. (Previously Presented) A compound or salt according to Claim 9, wherein

W is phenyl, which is unsubstituted or substituted with from 1 to 3 substituents independently selected from halogen, hydroxy, C₁₋₆alkoxy, -nitro, -CN, -SO₂NH₂, -SO₂NHR₆, -SO₂N(C₁₋₆alkyl)₂, amino, -NHC₁₋₆alkyl, -N(C₁₋₆alkyl)₂, -N(C₁₋₆alkyl)CO(C₁₋₆alkyl), -N(C₁₋₆alkyl)CO₂(C₁₋₆alkyl), -CONH₂, -CONH(C₁₋₆alkyl), -CON(C₁₋₆alkyl)₂, -CO₂(C₁₋₆alkyl), -S(C₁₋₆alkyl), -SO(C₁₋₆alkyl), -SO₂(C₁₋₆alkyl), and C₁₋₆alkyl optionally substituted with one or two substituents independently selected from hydroxy, halogen, and amino.

12. (Original) A compound or salt according to claim 9, wherein X is CH₂.

13. (Cancelled)

14. (Cancelled)

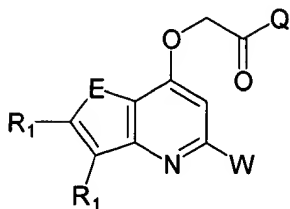
15. (Previously Presented) A compound or salt according to Claim 12; wherein

Q is pyridyl, which is unsubstituted or substituted with from 1 to 3 substituents independently selected from halogen, hydroxy, C₁₋₆alkoxy, -CN, amino, mono- and di(C₁₋₆)alkylamino, and C₁₋₆ alkyl which is unsubstituted or substituted with 1 or two substituents independently chosen from hydroxy, oxo, amino, halogen, C₁₋₆alkyl, C₁₋₆alkoxy, and mono- and di(C₁₋₆)alkylamino; and

W is phenyl which is unsubstituted or substituted with from 1 to 3 substituents independently selected from: halogen, hydroxy, C₁₋₆alkoxy, -nitro, -CN, -SO₂NH₂, -SO₂NHR₆, -SO₂N(C₁₋₆alkyl)₂, amino, -NHC₁₋₆alkyl, -N(C₁₋₆alkyl)₂, -N(C₁₋₆alkyl)CO(C₁₋₆alkyl), -N(C₁₋₆alkyl)CO₂(C₁₋₆alkyl), -CONH₂, -ONH(C₁₋₆alkyl), -CON(C₁₋₆alkyl)₂, -CO₂(C₁₋₆alkyl), -S(C₁₋₆alkyl), -SO(C₁₋₆alkyl), -SO₂(C₁₋₆alkyl), and C₁₋₆alkyl which is unsubstituted or substituted with one or two

substituents independently selected from hydroxy, halogen, and amino.

16. (Original) A compound or salt according to Claim 1 of formula:



17. (Cancelled)

18. (Previously Presented) A compound or salt according to Claim 16, wherein

W is phenyl which is unsubstituted or substituted with from 1 to 3 substituents independently selected from halogen, hydroxy, C₁₋₆alkoxy, -nitro, -CN, -SO₂NH₂, -SO₂NHR₆, -SO₂N(C₁₋₆alkyl)₂, amino, -NHC₁₋₆alkyl, -N(C₁₋₆alkyl)₂, -N(C₁₋₆alkyl)CO(C₁₋₆alkyl), -N(C₁₋₆alkyl)CO₂(C₁₋₆alkyl), -CONH₂, -ONH(C₁₋₆alkyl), -CON(C₁₋₆alkyl)₂, -CO₂(C₁₋₆alkyl), -S(C₁₋₆alkyl), -SO(C₁₋₆alkyl), -SO₂(C₁₋₆alkyl), and C₁₋₆alkyl which is unsubstituted or substituted with one or two substituents independently selected from hydroxy, halogen, and amino.

19. (Previously Presented) A compound or salt according to Claim 18, wherein:

Q is pyridyl, which is unsubstituted or substituted with from 1 to 3 substituents independently selected from: halogen, hydroxy, C₁₋₆alkoxy, -CN, amino, mono- and di(C₁₋₆)alkylamino, and C₁₋₆ alkyl which is unsubstituted or substituted with one or two substituents independently chosen from hydroxy, oxo, amino, halogen, C₁₋₆alkoxy, and mono- and di(C₁₋₆)alkylamino; and

W is phenyl which is unsubstituted or substituted with from 1 to 3 substituents independently selected from halogen, hydroxy, C₁₋₆alkoxy, -nitro, -CN, -SO₂NH₂, -SO₂NHR₆, -SO₂N(C₁₋₆alkyl)₂, amino, -NHC₁₋₆alkyl, -N(C₁₋₆alkyl)₂, -N(C₁₋₆alkyl)CO(C₁₋₆alkyl), -N(C₁₋₆alkyl)CO₂(C₁₋₆alkyl), -CONH₂, -CONH(C₁₋₆alkyl), -CON(C₁₋₆alkyl)₂, -CO₂(C₁₋₆alkyl), -S(C₁₋₆alkyl), -SO(C₁₋₆alkyl), -SO₂(C₁₋₆alkyl), and C₁₋₆alkyl which is unsubstituted or substituted with one or two substituents independently selected from hydroxy, halogen, and amino.

20-26. (Cancelled)

27. (Original) A compound according to Claim 1, which is 5-(4-Fluorophenyl)- 7-[(2-pyridyl)-methyloxy]-thieno[3,2-b]pyridine.

28. (Previously Presented) A compound according to Claim 1, which is 5-Phenyl-7-[(3-pyridyl)methyloxy]-thieno[3,2-b]pyridine.

29-32 (Cancelled)

33. (Previously Presented) A compound according to Claim 1, which is 7-[(4-Pyridyl)methyloxy]-5-phenylthieno[3,2-b]pyridine.

34-52. (Cancelled)

53. (Previously Presented) A pharmaceutical composition comprising a compound or salt according to Claim 1 combined with a pharmaceutically acceptable carrier or excipient.

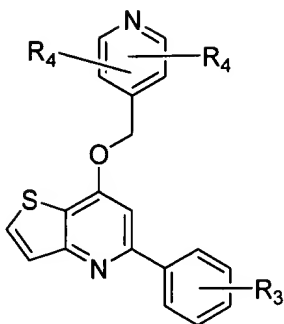
54-60. (Cancelled)

61. (Previously Presented) A method for the treatment of anxiety, depression, or a sleep disorder, comprising administering a therapeutically effective amount of a compound or salt of Claim 1 to a patient in need thereof.

62-66. (Canceled)

67-82 (Canceled)

83. (Previously Presented) A compound according to claim 1 of the formula



wherein

R₃ is selected from the group consisting of (C₁-C₆)alkyl, (C₁-C₆)alkoxy, halogen, and OH; and

R₄ at each occurrence is independently selected from the group consisting of hydrogen, halogen, hydroxy, alkoxy, -NO₂, -CN, -SO₂NH₂, -SO₂NH(C₁-C₆)alkyl, -SO₂N((C₁-C₆)alkyl)₂, amino, -NH(C₁-C₆)alkyl, -N((C₁-C₆)alkyl)₂, -N(R₆)CO((C₁-C₆)alkyl), -N((C₁-C₆)alkyl)CO₂((C₁-C₆)alkyl), -CONH₂, -CONH((C₁-C₆)alkyl), -CON((C₁-C₆)alkyl)₂, -CO₂((C₁-C₆)alkyl), and (C₁-C₆)alkyl.

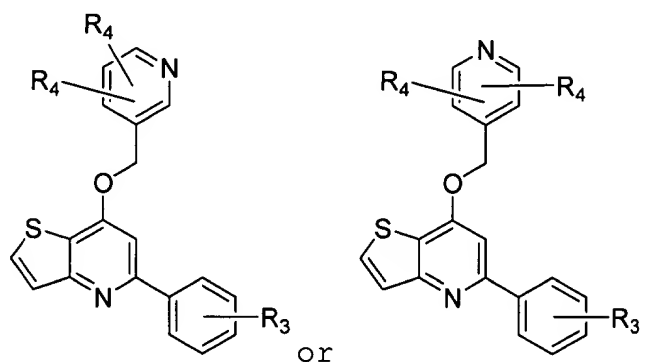
84. (Previously Presented) A compound according to claim 83, wherein

R₃ is selected from the group consisting of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, halogen, and OH; and
only one of the R₄ groups is hydrogen.

85. (Previously Presented) A compound according to claim 83, wherein

R₃ is selected from the group consisting of (C₁-C₄)alkyl, (C₁-C₄)alkoxy, halogen, and OH; and
one of the R₄ groups is halogen.

86. (Previously Presented) A compound according to claim 83, of the formula



87. (Previously Presented) A compound according to claim 86, wherein R₃ is halogen.